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European Patent Office
Office européen des brevets



(11) Publication number:

0 412 527 A2

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: 90115234.8

(51) Int. Cl.⁵: H01L 39/14, H01L 39/24

(22) Date of filing: 08.08.90

(30) Priority: 09.08.89 JP 206291/89
18.09.89 JP 241148/89

(43) Date of publication of application:
13.02.91 Bulletin 91/07

(84) Designated Contracting States:
DE FR GB IT

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(54) Ceramic superconductor wire and method of manufacturing the same.

(57) Disclosed herein is a ceramic superconductor comprising a ceramic superconductive member, and a high-conductivity metal layer (2) covering the ceramic superconductive member. At least one portion (3) of the metal layer having low electrical conductivity or low thermal conductivity. Also disclosed is a method of manufacturing a ceramic superconductor, comprising the steps of filling a ceramic superconductor or a precursor thereof in a high-conductivity metal pipe, thereby forming a composite member, rolling the composite member into a ceramic superconductor element of a desired shape, which comprises a ceramic superconductive member and a high-conductivity metal layer (2) covering the ceramic superconductive member, performing a predetermined heat treatment on the ceramic superconductor element, and alloying a predetermined portion of the high-conductivity metal layer (2), thereby rendering the portion less conductive either electrically or thermally.

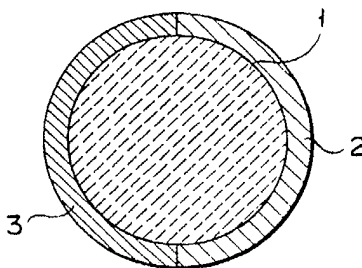


FIG. 1

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CERAMIC SUPERCONDUCTOR WIRE AND METHOD OF MANUFACTURING THE SAME

The present invention relates to a ceramic superconductor wire used as an electric wire, a cable, a coil winding, an electromagnetic shielding body, and a current lead wire for supplying a current to a superconductor element and a method of manufacturing the ceramic superconductor wire and, more particularly, a ceramic superconductor wire having a small AC current loss and a method of manufacturing the ceramic superconductor wire.

In recent years, various types of ceramic superconductors represented by chemical formulas such as $\text{YBa}_2\text{Cu}_3\text{O}_{7-y}$, $\text{Bi}_2\text{Sr}_2\text{Ca}_{n-1}\text{Cu}_n\text{O}_x$, $\text{Tl}_m\text{Ba}_2\text{Ca}_{n-1}\text{Cu}_n\text{O}_x$ ($m = 1, 2; n = 1, 2, 3, 4, 5, \dots$), and $(\text{Nd}_{1-x}\text{Ce}_x)_2\text{CuO}_{4-y}$ have been developed. Critical temperatures (T_c) of these ceramic superconductors reach 125 K.

Since these ceramic superconductors have poor workability, they are used in a state wherein a high-conductivity metal layer 2 is formed to cover a ceramic superconductor layer 1, as shown in Fig. 11. The form of such a ceramic superconductor is obtained by filling a ceramic superconductor or its precursor powder in a metal pipe or the like, rolling the pipe, and performing a predetermined heat treatment of the rolled pipe.

The high-conductivity metal layer consisting of a ceramic superconductor serves as a heat sink against heating caused by a flux jump as a rapid shift of a magnetic flux in the ceramic superconductor layer during energization. The high-conductivity metal layer also serves as a current bypath. A high-conductivity metal such as Ag or Cu having high thermal and electric conductivities is used as the above high-conductivity metal layer. In addition to these functions, a high-conductivity metal layer such as an Ag or Cu layer protects ceramic superconductors from external harmful substances and mechanically reinforces the ceramic superconductor.

A conventional ceramic superconductor wire covered with a high-conductivity metal layer such as an Ag or Cu layer causes the metal layer to generate an eddy current upon AC current energization of the ceramic superconductor wire.

The following problem is also posed when the above superconductor is used for a current lead. An SMES power generator, an MHD power generator, a fusion reactor, a linear motor car, a medical MRI, an accelerator magnet, and the like have been developed as low-temperature equipment using superconductor magnets. A current is supplied from an external power source to such low-temperature equipment through a current lead wire. The current lead wire consists of a metal material such as Cu, Cu-Ag, or Cu-P. Metal conductors are subjected to Joule heating upon current energization and absorption of a large amount of external heat. A large amount of coolant is undesirably evaporated.

There has been proposed use of a metal or intermetallic superconductor such as Nb-Ti and Nb_3Sn as the above current lead wire which does not produce Joule heat by cooling the metal or intermetallic superconductor to a liquid He temperature (4.2 K) to nullify an electric resistance. Such a superconductor cannot be used near any equipment except for low-temperature equipment cooled to the liquid He temperature and cannot sufficiently enhance its effect.

In most recent years, ceramic superconductors such as $\text{YBa}_2\text{Cu}_3\text{O}_7$, $\text{Bi}_2\text{Sr}_2\text{CaCu}_2\text{O}_8$, $\text{Bi}_2\text{Sr}_2\text{Ca}_2\text{Cu}_3\text{O}_{10}$, $\text{Tl}_2\text{Ba}_2\text{CaCu}_2\text{O}_8$, $\text{Tl}_2\text{Ba}_2\text{Ca}_2\text{Cu}_3\text{O}_{10}$ having critical temperatures (T_c) as high as 80 to 125 K, which allow substances to be superconductive at the liquid N_2 temperature, have been found.

These ceramic superconductors have high critical temperatures (T_c). When these superconductors are used as current lead wires, a liquid He cooling effect from the low-temperature equipment can be sufficiently utilized. Ceramic superconductors have low heat conductivities unlike metal materials such as Cu, and have an advantage in low absorption of external heat. Therefore, these ceramic superconductors have been expected as practical current lead wires.

Since a ceramic superconductor is, however, brittle, the following process is required. For example, a ceramic superconductor powder is filled in a metal tube as of Ag or Au, and the metal tube is then extruded, pressed, rolled, drawn or swaged. In this case, the metal tube must have good workability and a high purity, and have a ratio of a wall thickness to a sectional area to be 30 to 50% or more because the ceramic superconductor is hard. For this reason, a large amount of heat is conducted to the low-temperature equipment from a metal layer of a current lead wire rolled from the metal tube. A large amount of liquid He as a coolant is consumed, thus resulting in poor economy.

It is an object of the present invention to provide a ceramic superconductor wire having a high critical current density (J_c), a small AC current loss, and excellent productivity, and a method of manufacturing the ceramic superconductor wire.

It is another object of the present invention to provide a ceramic superconductor wire having a high constant critical current density (J_c) and a low heat conductivity and capable of minimizing an evaporation

amount of a coolant such as liquid He, and a method of manufacturing the ceramic superconductor wire.

The present invention is based on findings wherein part of a high-conductivity metal layer which covers a ceramic superconductor layer is converted into a low-conductivity portion, so that the AC current loss of the ceramic superconductor is greatly reduced. The present invention is also based on finding wherein a part of a high-conductivity metal layer which covers a ceramic superconductor layer is converted into a low-thermal conductivity portion, so that the evaporation amount of a coolant is greatly reduced. Extensive studies have also been made to reach the present invention.

According to the present invention, there is provided a ceramic superconductor comprising a ceramic superconductive member and a high-conductivity metal layer covering the ceramic superconductive member, at least a portion of the metal layer having low electrical conductivity or low thermal conductivity.

In the superconductor wire of the present invention, a low-conductivity portion is formed in the metal layer covered on the ceramic superconductor wire, and an eddy current loop generated by the metal layer during AC current energization is shielded by the low-conductivity portion, thereby reducing the AC current loss. Furthermore, in the present invention, a low thermal conductivity portion is formed in the metal layer covered on the ceramic superconductor wire, thereby reducing the evaporation amount of a coolant.

A metal such as Ag or Cu having high heat and electric conductivities is used to form the metal layer. An alloy obtained by solid-dissolving an element such as Pd, Ni, Zn, Cd, Sn, P, Si, or Be in the metal such as Ag or Cu is used to form the low-conductivity portion, which is formed in the metal layer. Same as the alloy obtained from Zn, In, Cd, Ar, Mg, Be, Ni, Fe, Co, Cr, Ti, Mn, Zr, Al, Ga, RE (Rare metal) in the metal such as Ag or Cu is used to form the low thermal conductivity portion.

The present invention will be described with reference to Figs. 1 to 5.

Referring to Fig. 1, the left half of a high-conductivity metal layer 2 which covers a ceramic superconductor layer 1 having a circular section is formed into a low-conductivity portion 3.

Fig. 2 shows a conductor tape. The lower half of a high-conductivity metal layer 2 is formed into a low-conductivity portion 3. This tape is suitable as an AC feeder cable conductor in a rotary structure. The tape is helically wound through an insulating film on a former (Al pipe) so that low-conductivity surfaces face each other. In this case, the former constitutes a 3-phase AC superconductor shown in Fig. 6. Reference numeral 21 in Fig. 6 denotes an Al pipe; 22, a superconductor tape; 23, semi-synthetic paper insulation; 24, a thermal insulation layer; and 25, a stainless steel pipe.

In a wire shown in Fig. 3, both ends of the conductor tape shown in Fig. 2 are slitted, and the slit portions are covered with resin layers 4. When this wire is wound around a core pipe serving as a coolant flow path, coupling losses produced between the wires can be appropriately minimized.

In a wire shown in Fig. 4, an outer half of a high-conductivity metal layer 2 of a ceramic superconductor layer 1 having a circular section is formed as a low-conductivity portion 3.

In a cable shown in Fig. 5, an entire metal layer of a ceramic superconductor layer 1 having a circular section is formed as a low-conductivity portion 3, and a plurality of the resultant wires are covered with a high-conductivity metal layer 2. The resultant cable is suitable as an AC magnet cable.

In the ceramic superconductor wires and cable shown in Figs. 1 to 5, each low-conductivity portion can be narrow. A better effect can be obtained when a plurality of narrow low-conductivity portions are formed at predetermined intervals so as to shield an eddy current loop.

According to the present invention, there is also provided a method of manufacturing a ceramic superconductor wire, comprising the steps of:
filling a ceramic superconductor or a precursor thereof in a high-conductivity metal pipe thereby forming a composite member;

rolling the composite member into a ceramic superconductor element of a desired shape, which comprises a ceramic superconductive member and a high-conductivity metal layer covering the ceramic superconductive member;

performing a predetermined heat treatment on the ceramic superconductor element; and

alloying a predetermined portion of the high-conductivity metal layer, thereby rendering said portion less conductive either electrically or thermally.

The following various ceramic superconductors may be used as a ceramic superconductor used in this method, such as $\text{YBa}_2\text{Cu}_3\text{O}_7$, $\text{Bi}_2\text{Sr}_2\text{CaCu}_2\text{O}_8$, $\text{Bi}_2\text{Sr}_2\text{Ca}_2\text{Cu}_3\text{O}_{10}$, $\text{Tl}_2\text{Ba}_2\text{CaCu}_2\text{O}_8$, $\text{Tl}_2\text{Ba}_2\text{Ca}_2\text{Cu}_3\text{O}_{10}$.

The precursor of the ceramic superconductor can be an intermediate produced in the process for synthesizing a ceramic superconductor from a starting material for a ceramic superconductor and can be exemplified as a mixture of oxides containing elements constituting a ceramic superconductor, a co-precipitated mixture, an oxygen-deficient compound oxide, or an alloy constituted by the above alloy elements. Such a precursor is reacted with a ceramic superconductor during heating in an oxygen atmosphere.

When the ceramic superconductor is an oxygen-deficient n-type oxide superconductor, a material of the high-conductivity metal pipe can be Cu which can be easily oxidized in a reduction atmosphere. However, when the ceramic superconductor consists of a p-type oxide superconductor such as an oxide superconductor represented by chemical formula $\text{YBa}_2\text{Cu}_3\text{O}_{7-\delta}$ (δ is advantageously reduced to almost zero), heating is performed in the oxygen atmosphere. A material of the metal pipe is preferably Ag which cannot be easily oxidized and has a high O_2 permeability. Another metal such as Au, Pd, Pt, Ir, Rh is preferable in consideration of non-reactivity with a superconductor and mechanical workability.

A method of forming a low-conductivity portion in a high-conductivity metal layer may be: a method of precipitating an element for reducing an electric conductivity at a predetermined portion of the surface of the high-conductivity metal layer in accordance with plating, PVD (Physical Vapor Deposition), or CVD (Chemical Vapor Deposition), thermally diffusing the precipitated element, and alloying the precipitated element with the high-conductivity metal layer to form a low-conductivity portion; ion implantation; or a method obtained by combining ion implantation and thermal diffusion.

According to the method of the present invention, Assume that the ceramic superconductor is a p-type oxide superconductor and alloying is performed in the high-conductivity metal layer to form a low-conductivity portion prior to heating of a rolled pipe. In this case, when an alloy element is a base element, the alloy element is oxidized to restore the electrical conductivity of the low-conductivity portion. Therefore, mixing of the alloy element is preferably performed after heating.

A method of forming a low-thermal conductivity portion can be performed as follows. A film is formed on a metal layer by exotic metal plating, PVD, CVD, paste coating, soldering, cladding, or the like and is heated to diffuse an exotic metal in the metal layer to alloy the exotic metal with a metal constituting the metal layer. Examples of the exotic metal are Zn, In, Cd, Cu, Mg, Be, Ni, Fe, Co, Cr, Ti, Mn, Zr, Al, Ga, and RE (rare-earth element). Such a metal is suitable because it can greatly reduce the heat conductivity of the metal layer in a small amount.

Another useful alloying method is ion implantation or a method of holding a ceramic superconductor wire in a metal vapor to partially alloy the wire with the metal vapor. A terminal connected to low-temperature equipment or an external power source is desirably kept to have a low electric resistance without requiring alloying so as to suppress Joule heating.

This method will be described with reference to Figs. 7 to 10.

Figs. 7 and 8 are perspective views of ceramic superconductor wires obtained by covering the surfaces of ceramic superconductors with metal layers, respectively.

More specifically, the ceramic superconductor wire shown in Fig. 7 is obtained by covering a rod-like ceramic superconductor powder-molded body 11 with a metal layer 2. The ceramic superconductor wire shown in Fig. 8 is obtained by covering noble metal layers 12 on the upper and lower surfaces of a plate-like ceramic superconductor powder-molded body 11. End portions of the ceramic superconductor powder-molded body 1 are exposed outside.

A ceramic superconductor wire is heated and sintered at a predetermined temperature to partially alloy the metal layer 2 so as to form portions 13 except for connecting portions 14 of the ceramic superconductor, as shown in Fig. 9.

As shown in Fig. 10, an improved current lead wire is obtained such that noble metal layers 12 of the portions 13 of the ceramic superconductor wire except for the connecting portions 14 are made thin. The thin noble metal layers 12 can further reduce an amount of absorption heat. In this case, the thin noble metal layers 12 are formed, and then connecting portions are formed at terminals of the wire. Alloying can be performed before or after the thin noble metal layers 12 are formed. A method of forming thin noble metal layers may be any method such as chemical etching or mechanical milling.

A single-core superconductor wire has been exemplified. The method of the present invention is, however, equally applicable to a multi-core superconductor cable having a plurality of superconductor wires.

The ceramic superconductor wire according to the present invention has the low-conductivity portion partially formed in the high-conductivity metal layer as the outermost layer of the ceramic superconductor wire. Generation of an eddy current during AC current energization can be suppressed, and the AC current loss can be greatly reduced. Since the low-conductivity metal portion may be formed in part of the high-conductivity metal layer, the main function of the high-conductivity metal layer as a heat sink against a flux jump cannot be impaired. In addition, the function of protecting the ceramic superconductor layer against the outer atmosphere is not lost from the metal layer. Moreover, since the low-conductivity portion is formed by alloying, mechanical properties of the metal layer can be improved.

Positive conductivity correlation between heat and electricity is apparently established in metals without citing the Wiedemann-Franz law. Wire materials of the present invention are suitable as current lead wires.

According to the present invention, in a high- T_c ceramic superconductor whose at least part of the

surface is covered with a high-conductivity layer as of a noble metal, the noble metal layer is alloyed to form a current lead wire. In use of the resultant current lead wire, Joule heating of the ceramic superconductor wire can be suppressed by a cooling effect of the low-temperature equipment, and conduction of external heat to the low-temperature equipment due to a decrease in heat conductivity of the ceramic superconductor wire can be suppressed.

Alloying of the noble metal layer is performed after predetermined rolling of the ceramic superconductor wire finishes. Therefore, rolling workability of the ceramic superconductor wire is not impaired. The noble metal layer is made thin before or after the noble metal is alloyed, thereby further suppressing heat conduction from the current lead wire.

This invention can be more fully understood from the following detailed description when taken in conjunction with the accompanying drawings, in which:

Fig. 1 is a view showing a ceramic superconductor wire using a ceramic superconductor layer having a circular section according to the present invention;

Fig. 2 is a view for explaining a ceramic superconductor wire employing a conductor tape according to the present invention;

Fig. 3 is a view for explaining a ceramic superconductor wire having conductive tapes respectively formed at both ends of the wire according to the present invention;

Fig. 4 is a view for explaining a ceramic superconductor wire having a high-conductivity metal layer whose outer half is formed into a low-conductivity portion according to the present invention;

Fig. 5 is a view for explaining a ceramic superconductor wire having a metal layer which is entirely formed as a low-conductivity portion according to the present invention;

Fig. 6 is a view for explaining a 3-phase AC superconductor cable employing a ceramic superconductor wire;

Figs. 7 and 8 are perspective views of a ceramic superconductor wire obtained by covering a ceramic superconductor wire with a metal layer according to the present invention;

Fig. 9 is a view for explaining an alloyed ceramic superconductor wire according to the present invention;

Fig. 10 is a view for explaining a ceramic superconductor wire whose thin metal layer is alloyed according to the present invention; and

Fig. 11 is a view showing a sectional structure of a conventional wire.

The present invention will be described in detail by way of its examples.

Example 1

Y_2O_3 , $BaCO_3$, and CuO were blended and mixed at an atomic ratio of 1 : 2 : 3 (Y : Ba : Cu), and a powder mixture was calcined in air at $880^\circ C$ for 24 Hr. The calcined body was pulverized and classified to obtain a calcined powder. The calcined powder was molded by CIP molding to form a 9-mm diameter rod. The rod was preliminarily sintered in an oxygen flow at $850^\circ C$ for 8 Hr and was filled in an Ag pipe having an outer diameter of 12 mm and an inner diameter of 9 mm. The Ag pipe filled with the CIP-molded body was swaged and rolled by a grooved roll to obtain a 2.5-mm wire. This wire was formed into a 0.3-mm thick tape by a roll. The tape element was heated in an oxygen flow at $900^\circ C$ for 8 Hr and was then cooled from $900^\circ C$ at a rate of $2^\circ C/min$. Zn was electrically plated on one surface of the tape element to a thickness of $4.5 \mu m$, and the resultant structure was heated in an Ar flow at $450^\circ C$ for 10 Hr. This tape was slitted to have a width of 3 mm, and a Teflon tape was wound around each wire to obtain a ceramic superconductor wire.

Example 2

Bi_2O_3 , SrO , CaO , and CuO were blended and mixed at an atomic ratio of 2 : 1.9 : 2.1 : 2.5 (Bi : Sr : Ca : Cu), and a powder mixture was calcined in an oxygen flow at $650^\circ C$. The calcined body was pulverized and classified to obtain a calcined powder. Following the same procedures as in Example 1, the calcined powder was subjected to CIP molding, sealing in an Ag pipe, and rolling to form a 0.3-mm thick tape element.

The Ag layer on one surface of the tape element was etched with HNO_3 by a depth of $30 \mu m$, Pd was plated on the etched surface by a thickness of $10 \mu m$. The resultant structure was held in an $N_2 + 7.5\%O_2$ flow at $890^\circ C$ for 30 minutes and was cooled from $890^\circ C$ at a rate of $2.5^\circ C/min$. Both ends of the resultant structure were slitted to have a width of 3 mm, and a Teflon tape was wound around each wire for electric

insulation, thereby obtaining a ceramic superconductor wire.

Example 3

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Nd_2O_3 , CeO_2 , and CuO were blended and mixed at an atomic ratio of 1.7 : 0.3 : 1 (Nd : Ce : Cu), and a powder mixture was calcined in an O_2 flow at $1,050^\circ\text{C}$ for 8 Hr. The calcined body was pulverized and classified to obtain a calcined powder. The calcined powder was CIP-molded to obtain a 9-mm diameter rod. The rod was filled in an Ni-plated Cu pipe having an outer diameter of 12 mm and an inner diameter of 9 mm after the rod was preliminarily sintered.

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Following the same procedures as in Example 1, the Cu pipe filled with the preliminarily sintered rod body was rolled into a 0.3-mm thick tape element. Zn and Ni were sequentially plated on one surface of the tape element to thicknesses of $5\text{ }\mu\text{m}$ and $10\text{ }\mu\text{m}$, respectively. The resultant structure was heated in an Ar flow at 980°C for 4 Hr and was removed from a heating furnace and cooled. Both ends of the sintered body were slitted into a tape body having a width of 3 mm. A Teflon tape was wound around each wire, thereby obtaining a ceramic superconductor wire.

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Critical current densities (J_c) and conductivities of the metal layers of the ceramic superconductor wires prepared in Examples 1 to 3 were measured. Test results are shown in Table 1 below.

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As is apparent from Table 1, the samples in Examples 1 to 3 had high critical current densities (J_c). The conductivities of the low-conductivity portions partially formed in the high-conductivity metal layers which cover the ceramic superconductors were as low as 1 to 4%. The eddy currents generated in the metal layers during AC current energization can be greatly suppressed. For this reason, the AC current loss can be reduced to about 10% of the conventional wire whose metal layer is a rigid conductive metal layer.

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The samples in Examples 1 to 3 exemplified wires having the structure shown in Fig. 2 or 3. The entire metal layers of the wires may be alloyed, and a plurality of wires are bundled, soldered, and fixed within an Ag or Cu pipe to obtain a cable shown in Fig. 5. This also provides the same effect as in Examples 1 to 3.

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Table 1

Classification	No.	Type of Super-conductor	Metal Layer		J_C^* A/cm ²	Conduc-tivity Ratio ^{**}
			High-Conductivity Metal Layer	Low-Conductivity Portion		
Sample of Present Invention	Example 1	Y	Ag	Zn mixing	0.8×10^4	2.1
	Example 2	Bi	Ag	Pd mixing	5.1×10^4	3.5
	Example 3	Nd	Cr-plated Cu	Zn, Ni mixing	1.2×10^4	1.3

* Measured in a magnetic field of 500 gauss at 40 K for sample Nos. 1 and 2.

Measured in a magnetic field of 500 gauss at 4.2 K for sample No. 3.

** The ratio of conductivity of low-conductivity portion to that of high-conductivity metal layer

Example 4

5 Bi_2O_3 , SrO , CaO , CuO powders were blended and mixed at an atomic ratio of 2 : 2 : 1 : 2 (Bi : Sr : Ca : Cu) and this powder mixture was calcined in air at 800°C for 30 Hr. The calcined body was pulverized and classified to obtain a $\text{Bi}_2\text{Sr}_2\text{CaCu}_2\text{O}_x$ calcined powder. This calcined powder was filled in an Ag tube having an outer diameter of 26 mm and an inner diameter of 18 mm. The tube with the calcined powder is rolled
10 by a grooved roll to obtain a 4-mm diameter wire. This wire was rolled by flat rolling to obtain a tape having a thickness of 0.3 mm and a width of 7 mm.

The tape was sintered in an O_2 flow at 860°C for 30 Hr to obtain an oxide superconductor wire covered with an Ag layer. The Ag layer of the oxide superconductor wire was plated with Zn to form a $3\text{-}\mu\text{m}$ thick Zn layer. The resultant structure was heated in an Ar flow at 450°C for 2 Hr to prepare a current lead wire.

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Example 5

A current lead wire was manufactured following the same procedures as in Example 4 except that the
20 thickness of Zn plated on the Ag layer was $10\text{ }\mu\text{m}$ and heating was performed at 450°C for 4 Hr.

Example 6

25 A current lead wire was manufactured following the same procedures as in Example 4 except that Zn was plated on an Ag layer to a thickness of $3\text{ }\mu\text{m}$, Cd was then plated to have a thickness of $5\text{ }\mu\text{m}$, and heating was performed at 500°C for 2 Hr.

30 Examples 7 - 9

Current lead wires were respectively manufactured following the same procedures as in Examples 4 to 6 except that surfaces of Ag layers of oxide superconductor wires covered with the Ag layers were etched by a diluted nitric acid by a depth of $15\text{ }\mu\text{m}$ each.

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Comparative Examples 1 and 2

Current lead wires were manufactured following the same procedures as in Example 4 or 7 except that
40 Ag layers were not alloyed.

Jc values and heat conductivities of the current lead wires of Examples 4 to 9 and Comparative Examples 1 and 2 were measured at 4.2 K and 77 K. Test results are summarized together with main manufacturing conditions in Table 2 below.

As is apparent from Table 2, samples (No. 4 to 9) of the present invention had lower heat conductivities
45 than those of samples (No. 10 and 11) of the comparative examples due to alloying of the metal layers on the surfaces of the ceramic superconductor wires.

In the current lead wires of sample Nos. 7 to 9, the Ag layers on the ceramic superconductor wires were etched and made thin, and thin Ag layers were alloyed. Therefore, the alloying density was increased to further reduce the heat conductivities. Therefore, external heat conduction could be greatly suppressed
50 together with an effect of thin Ag layers.

The wires of sample Nos. 4 to 9 had lower critical current densities (Jc) than those of sample Nos. 10 and 11 of the comparative examples by amounts corresponding to the degrees of alloying of the Ag layers formed on the ceramic superconductor wires, but differences were very small.

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Table 2

	Classification	No.	Surface Treatment of Ag Film	Alloying		4.2 K		77 K	
				Plating Thickness (μm) Zn + Cd	Heating Condition $^{\circ}\text{C} \times \text{H}$	J_c A/cm ²	Heat Conductivity*	J_c A/cm ²	Heat Conductivity*
Sample by Method of Present Invention	Example 4	4	No	3	450 \times 2	3.9×10^5	38	1.7×10^4	51
	Example 5	5	No	10	450 \times 4	3.7×10^5	30	1.4×10^4	47
	Example 6	6	No	3 + 5	500 \times 2	3.7×10^5	32	1.5×10^4	49
	Example 7	7	Yes	3	450 \times 2	3.6×10^5	30	1.5×10^4	42
		8	Yes	10	450 \times 4	3.5×10^5	27	1.2×10^4	37
		9	Yes	3 + 5	500 \times 2	3.4×10^5	28	1.4×10^4	37
	Comparative Example 1	10	No	-	-	3.9×10^5	100	2.0×10^4	100
	Comparative Example 2	11	Yes	-	-	3.8×10^5	78	1.9×10^4	79
	Sample by Comparative Method								

* Relative values with respect to Comparative Examples (Nos. 10 and 11) as 100

Claims

- 5 1. A ceramic superconductor comprising:
a ceramic superconductive member; and
a high-conductivity metal layer (2) covering the ceramic superconductive member, at least a portion (3) of
said metal layer having low electrical conductivity or low thermal conductivity.
- 10 2. A ceramic superconductor according to claim 1, characterized in that that portion of said metal layer
which has the low electrical conductivity is made of an alloy containing at least one element selected from
the group consisting of Pd, Ni, Zn, Cd, Sn, P, Si, and Be.
3. A ceramic superconductor according to claim 1, characterized in that that portion of said metal layer
which has the low thermal conductivity is made of an alloy containing at least one element selected from
15 the group consisting of Zn, In, Cd, Cu, Mg, Be, Ni, Fe, Co, Cr, Ti, Mn, Zr, Al, Ga, and a rare-earth element.
4. A ceramic superconductor according to claim 1, characterized in that the remaining portion (3) of said
metal layer (2) which has high electrical conductivity is made of one material selected from the group
consisting of Cu, a Cu alloy, Ag, an Ag alloy, and a heat-resistant nickel alloy.
5. A ceramic superconductor according to claim 1, characterized in that the remaining portion of said metal
20 layer (2) which has high electrical conductivity is made of a noble metal selected from the group consisting
of Ag, Au, Pd, Pt, Ir, and Rh.
6. A ceramic superconductor according to claim 1, characterized in that said ceramic member is made of a
compound selected from the group consisting $\text{YBa}_2\text{Cu}_3\text{O}_7$, $\text{Bi}_2\text{Sr}_2\text{CaCu}_2\text{O}_8$, $\text{BiSr}_2\text{Ca}_2\text{Cu}_3\text{O}_{10}$,
 $\text{Tl}_2\text{Ba}_2\text{CaCu}_2\text{O}_8$, and $\text{Tl}_2\text{Ba}_2\text{Ca}_2\text{Cu}_3\text{O}_{10}$.
- 25 7. A ceramic superconductor according to claim 1, characterized in that said ceramic superconductive
member is a wire having a circular cross section.
8. A ceramic superconductor according to claim 1, characterized in that said ceramic superconductive
member is a tape having a rectangular cross section.
9. A ceramic superconductor according to claim 1, characterized in that said ceramic superconductive
30 member comprises a bundle of ceramic wires each covered by a metal layer having low electrical
conductivity, and a metal layer covering the bundle of ceramic wires and having high electrical conductivity.
10. A ceramic superconductor according to claim 1, characterized in that said ceramic superconductive
member is a current-guiding conductor which has low thermal conductivity, except for both end portions
thereof.
- 35 11. A method of manufacturing a ceramic superconductor, characterized by comprising the steps of:
filling a ceramic superconductor or a precursor thereof in a high-conductivity metal pipe, thereby forming a
composite member;
rolling the composite member into a ceramic superconductor element of a desired shape, which comprises
a ceramic superconductive member and a high-conductivity metal layer covering the ceramic superconduc-
40 tive member;
performing a predetermined heat treatment on the ceramic superconductor element; and
alloying a predetermined portion (2) of the high-conductivity metal layer, thereby rendering said portion less
conductive either electrically or thermally.
12. A method according to claim 11, characterized in that the step of alloying the predetermined portion of
45 the high-conductivity metal layer (2) comprises the step of forming a layer of an alloying metal on the
predetermined portion of said high-conductivity metal layer by means of plating, PVD, CVD, paste-coating,
soldering, cladding, or thin-film forming, and the step of thermally diffusing the alloying metal into the
predetermined portion of said high-conductivity metal layer (2).
13. A method according to claim 11, characterized in that the step of alloying the predetermined portion of
50 said high-conductivity metal layer (2) is either the step of injecting ions of an alloying metal into the
predetermined portion of said high-conductivity metal layer (2) or the step of both injecting ions of an
alloying metal and diffusing the alloying metal into the predetermined portion of said high-conductivity metal
layer (2).
14. A method according to claim 11, characterized in that said step of alloying the predetermined portion of
55 the high-conductivity metal layer (2) is the step of placing the predetermined portion of said high-
conductivity metal layer (2) in vapor of the alloying metal.
15. A method according to claim 11, characterized in that said high-conductivity metal pipe is made of one
material selected from the group consisting of Cu, a Cu alloy, Ag, an Ag alloy, and a heat-resistant nickel

alloy.

16. A method according to claim 12, characterized in that said alloying metal is made of at least one element selected from the group consisting Zn, In, Cd, Cu, Mg, Be, Ni, Fe, Co, Cr, Ti, Mn, Zr, Al, Ga, and a rare-earth element.

5 17. A method according to claim 12, characterized by further comprising the step of making the predetermined portion of said high-conductivity metal layer (2) thinner prior to said step of alloying the predetermined portion of the high-conductivity metal layer.

18. A method according to claim 11, characterized in that said ceramic superconductor which is filled in said high-conductivity metal pipe is one selected from the group consisting of $\text{YBa}_2\text{Cu}_3\text{O}_7$, $\text{Bi}_2\text{Sr}_2\text{CaCu}_2\text{O}_8$,
10 $\text{BiSr}_2\text{Ca}_2\text{Cu}_3\text{O}_{10}$, $\text{Tl}_2\text{Ba}_2\text{CaCu}_2\text{O}_8$, and $\text{Tl}_2\text{Ba}_2\text{Ca}_2\text{Cu}_3\text{O}_{10}$.

19. A method according to claim 11, characterized in that the precursor of said ceramic superconductor, which is filled in said high-conductivity metal pipe, is one selected from the group consisting of oxides of elements forming an intermediate ceramic superconductor, coprecipitate of two or more of said elements, oxygen-depleted composite oxide of two or more of said elements, and alloys of two or more of said
15 elements.

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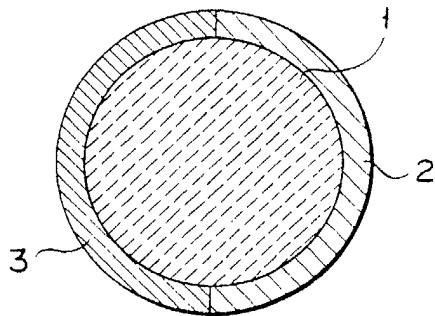


FIG. 1

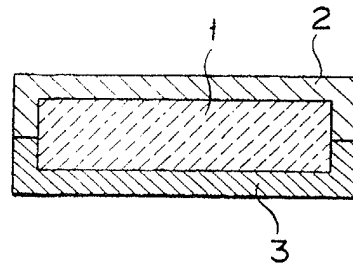


FIG. 2

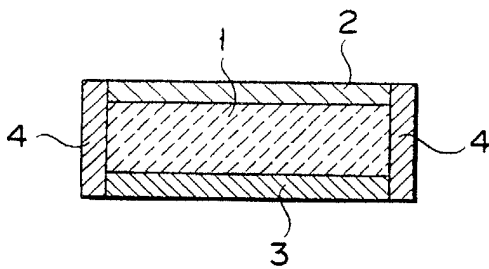


FIG. 3

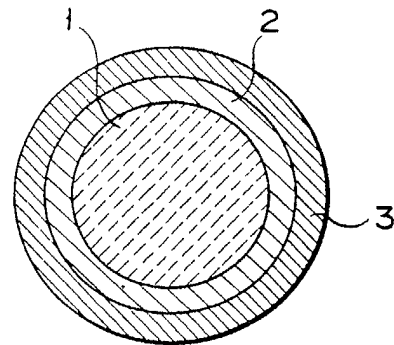


FIG. 4

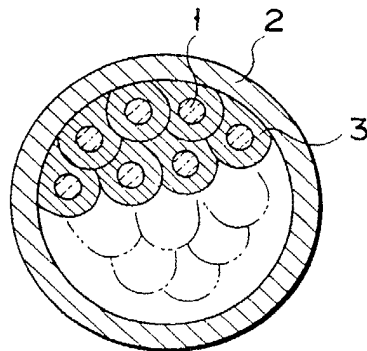
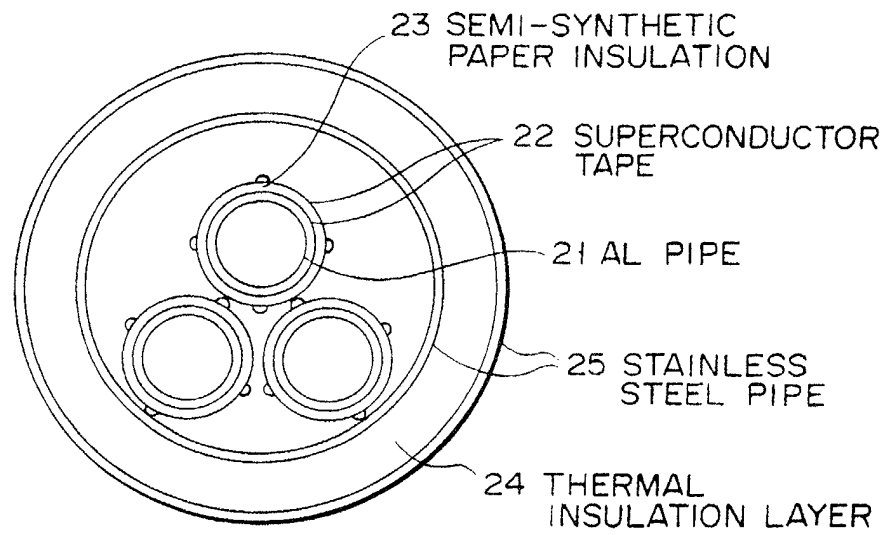
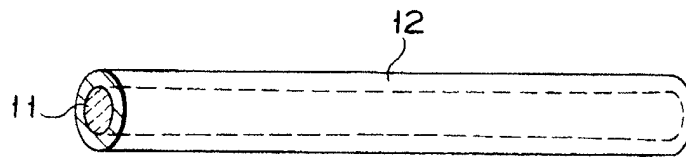


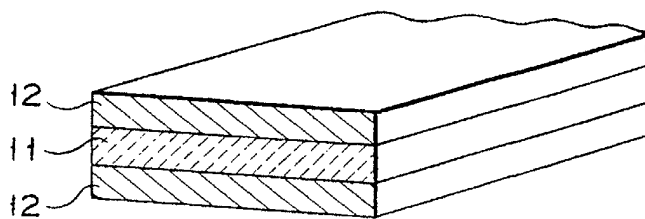
FIG. 5



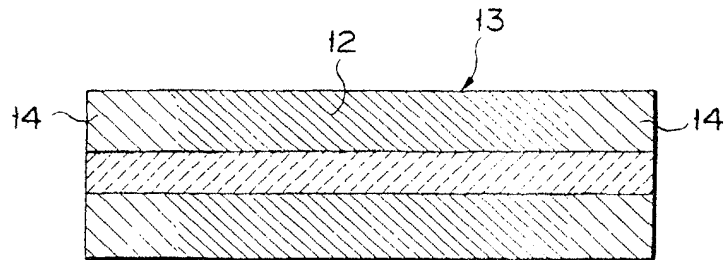
F I G. 6



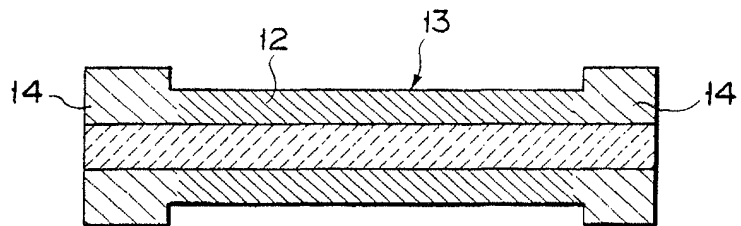
F I G. 7



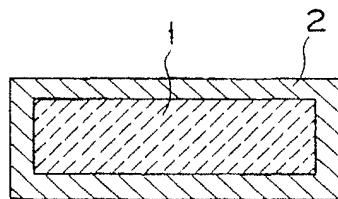
F I G. 8



F I G. 9



F I G. 10



F I G. 11